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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
10/758,683	01/14/2004	Jeannette Whitcomb	. 11068-078-999	4994
7590 11/28/2007 JONES DAY			EXAMINER	
222 East 41st Street			PARKIN, JEFFREY S	
New York, NY 10017			ART UNIT	PAPER NUMBER
			1648	
·			MAIL DATE	DELIVERY MODE
			11/28/2007	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)			
·	10/758,683	WHITCOMB, JEANNETTE			
Office Action Summary	Examiner	Art Unit			
	Jeffrey S. Parkin, Ph.D.	1648			
The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REF WHICHEVER IS LONGER, FROM THE MAILING  - Extensions of time may be available under the provisions of 37 CFR after SIX (6) MONTHS from the mailing date of this communication.  - If NO period for reply is specified above, the maximum statutory period.  - Failure to reply within the set or extended period for reply will, by state Any reply received by the Office later than three months after the ma earned patent term adjustment. See 37 CFR 1.704(b).	DATE OF THIS COMMUNIC 1.136(a). In no event, however, may a re od will apply and will expire SIX (6) MONT tute, cause the application to become ABA	ATION. ply be timely filed  HS from the mailing date of this communication. ANDONED (35 U.S.C. § 133).			
Status		•			
1) Responsive to communication(s) filed on 12	September 2007.				
2a) ☐ This action is <b>FINAL</b> . 2b) ☑ TI	This action is <b>FINAL</b> . 2b)⊠ This action is non-final.				
3) Since this application is in condition for allow	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is				
closed in accordance with the practice unde	r Ex parte Quayle, 1935 C.D.	11, 453 O.G. 213.			
Disposition of Claims		•			
4) ☐ Claim(s) 83-89 is/are pending in the applicate 4a) Of the above claim(s) is/are withd 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) 83-89 is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) are subject to restriction and	rawn from consideration.				
Application Papers					
9) The specification is objected to by the Exami 10) The drawing(s) filed on is/are: a) a Applicant may not request that any objection to the Replacement drawing sheet(s) including the correction.  The oath or declaration is objected to by the	ccepted or b) objected to be drawing(s) be held in abeyand ection is required if the drawing(s	ce. See 37 CFR 1.85(a). s) is objected to. See 37 CFR 1.121(d).			
Priority under 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of:  1. Certified copies of the priority docume 2. Certified copies of the priority docume 3. Copies of the certified copies of the priority docume application from the International Bure * See the attached detailed Office action for a life.	ents have been received. ents have been received in Apriority documents have been reau (PCT Rule 17.2(a)).	oplication No received in this National Stage			
Attachment(s)	_				
<ol> <li>Notice of References Cited (PTO-892)</li> <li>Notice of Draftsperson's Patent Drawing Review (PTO-948)</li> <li>Information Disclosure Statement(s) (PTO/SB/08)</li> <li>Paper No(s)/Mail Date</li> </ol>	Paper No(s)	ummary (PTO-413) /Mail Date formal Patent Application _			

Serial No.: 10/758,683 Docket No.: 11068-078-999
Applicant: Whitcomb, J. Filing Date: 01/14/2004

#### Detailed Office Action

#### Status of the Claims

Acknowledgement is hereby made of receipt and entry of the communication filed 12 September, 2007. Claims 83-89 are pending in the instant application.

## 35 U.S.C. § 102

The following is a quotation of the appropriate paragraphs of 35 U.S.C. § 102 that form the basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless --

(a) the invention was known or used by others in this country, or patented or described in a printed publication in this or a foreign country, before the invention thereof by the applicant for a patent.

The previous rejection of claims 83-85 and 89 under 35 U.S.C. § 102(a) as being clearly anticipated by Fujiwara et al. (1998), is hereby withdrawn in response to applicant's amendment.

## 35 U.S.C. § 112, First Paragraph

The following is a quotation of the first paragraph of 35 U.S.C. § 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

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## Scope of Enablement

Claims 83-89 are rejected under 35 U.S.C. § 112, first paragraph, because the specification does not reasonably enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the invention commensurate in scope with these claims. The claims are directed toward a method for assessing the effectiveness of NNRTI on an HIV-infected patient comprising evaluating nucleic acid samples obtained from patient plasma for certain Appropriately drafted claim language mutations in the RT. directed toward HIV-1-infected patients and the mutations identified in the specification would be appropriate Subject Matter below for suggested claim (see **Allowable** language).

considerations enablement The legal that govern determinations pertaining to undue experimentation have been clearly set forth. Enzo Biochem, Inc., 52 U.S.P.Q.2d 1129 In re Wands, 8 U.S.P.Q.2d 1400 (C.A.F.C. (C.A.F.C. 1999). 1988). Ex parte Forman 230 U.S.P.Q. 546 (PTO Bd. Pat. App. courts concluded that several Int., 1986). The factual inquiries should be considered when making such assessments including the quantity of experimentation necessary, the amount of direction or guidance presented, the presence or absence of working examples, the nature of the invention, the state of the prior art, the relative skill of those in that art, the predictability or unpredictability of the art and the breadth of the claims. In re Rainer, 52 C.C.P.A. 1593, 347 F.2d 574, 146 U.S.P.Q. 218 (1965). The disclosure fails to provide adequate guidance pertaining to a number of these considerations as follows:

The disclosure fails to provide adequate guidance pertaining to the genotypic/phenotypic properties of other RTs as it pertains to NNRTI increased/decreased susceptibility. The NNRTIs interact in a highly specific manner with the RT based upon steric constraints. Thus, only a limited number of amino acid mutations are permitted that lead to the desired phenotype. Thus, the skilled artisan would readily question the ability of other mutations at the disclosed location to produce a mutant RT with the desired phenotype.

The disclosure fails to provide sufficient guidance pertaining to the presence of the claimed mutations in the human immunodeficiency virus type 2 (HIV-2) RT. Although HIV-1 and -2 are both lentiviruses, they only display ~35-38% genetic relatedness at the nucleotide sequence level. Accordingly, considering the genetic unrelatedness between these two viruses, it seems improbable that the same mutations in HIV-1 RT would be present in the HIV-2 RT.

The prior art teaches that a limited number of substitutions are associated with increased/decreased NNRTI drug resistance (Fujiwara et al., 1998; Bacheler et al., 2001; Ceccherini-Silberstein et al., 2007). However, the claims are broadly directed toward any combination of amino acid substitutions. This constitutes a rather large genus of variants which are not supported by the disclosure. For instance, would the skilled artisan reasonably expect G190W and K101H to display increased/decreased susceptibility to NNRTIs?

Accordingly, when all the aforementioned factors are considered *in toto*, it would clearly require undue experimentation to practice the claimed invention in a manner commensurate in scope with the claims.

#### Allowable Subject Matter

As previously set forth, it appears that the combination of mutations G190A/S, K101E, and A98G are free of the prior art. Appropriately drafted claim language directed toward these combinations of mutations would be allowable (i.e., A method of assessing the effectiveness of a nonnucleoside transcriptase inhibitor (NNRTI) on human immunodeficiency a virus type 1 (HIV-1)-infected patient, comprising evaluating whether a plasma sample colected from the HIV-1-infected patient comprises a encoding a reverse transcriptase (RT) that has one of the following mutations: G190A/S and K101E; G190A/S, K101E, and K103N; G190A/S and A98G; (etc.), wherein said mutation is associated with an increased susceptiblity to...).

# Correspondence

Any inquiry concerning this communication should be directed to Jeffrey S. Parkin, Ph.D., whose telephone number is (571) 272-0908. The examiner can normally be reached Monday through Thursday from 10:30 AM to 9:00 PM. A message may be left on the examiner's voice mail service. If attempts to reach the examiner are unsuccessful, the examiner's supervisor, Bruce R. Campell, Ph.D., can be reached at (571) 272-0974. Direct general status inquiries to the Technology Center 1600 receptionist at (571) 272-1600. Informal communications may be submitted to the Examiner's RightFAX account at (571) 273-0908.

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Updated Notice of Centralized Delivery and Facsimile Transmission Policy for Patent Related Correspondence, and Exceptions Thereto, 1292 Off. Gaz. Pat. Office 186 (March 29, 2005).

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Respectfully,

Jeffrey S. Parkin, Ph.D.

Primary Examiner Art Unit 1648

26 November, 2007